IN THE CLAIMS

Please amend claims 38 and 67 and cancel claims 61-65 without any prejudice or disclaimer to the subject matter expressed therein as indicated in the complete listing of all claims in the application set forth below.

Claims 1-37 (canceled)

Claim 38 (currently amended): A method of improving patient compliance with a therapeutic or nutritional regimen, which comprises:

administering to an animal a non-effervescent flavored suspension formed by placing into a liquid a solid dispersible tastemasked tablet comprising a flavoring agent and a plurality of particles being coated with an extended release coating agent;

wherein the solid dispersible tablet forms a non-effervescent flavored suspension when placed in a liquid having a viscosity of about 25 cp to about 75 cp; and

wherein the non-effervescent flavored suspension after being orally administered to the animal releases the biologically active substance over a period of about 2 hours to about 48 hours.

Claim 39 (original): The method of claim 38, wherein the non-

effervescent flavored suspension is formed in less than about 10 minutes after the solid dispersible tablet is placed in the liquid.

Claim 40 (original): The method of claim 38, wherein the noneffervescent flavored suspension is formed in less than about 5 minutes after the solid dispersible tablet is placed in the liquid.

Claim 41 (original): The method of claim 38, wherein the noneffervescent flavored suspension is formed in less than about 1 minute after the solid dispersible tablet is placed in the liquid.

Claim 42 (original): The method of claim 38, wherein the noneffervescent flavored suspension is formed in less than about 30 seconds after the solid dispersible tablet is placed in the liquid.

Claim 43 (original): The method of claim 38, wherein the noneffervescent flavored suspension is formed upon stirring, mixing or blending the liquid after the solid dispersible tablet is placed in said liquid.

Claim 44 (original): The method of claim 38, wherein the noneffervescent flavored suspension is formed without stirring, mixing or blending the liquid after the solid dispersible tablet is placed in said liquid.

Claim 45 (previously presented): The method of claim 38, wherein the solid dispersible tablet is a self-dispersing tablet.

Claim 46 (original): The method of claim 38, wherein the non-effervescent flavored suspension after being orally administered to the animal releases the biologically active substance for a period of about 4 hours up to about 24 hours.

Claim 47 (original): The method of claim 38, wherein the non-effervescent flavored suspension after being orally administered to the animal releases the biologically active substance for a period of about 12 hours up to about 24 hours.

Claim 48 (original): The method of claim 38, wherein the solid dispersible tablet further contains a coloring agent, and wherein the suspension is a colored suspension.

Claim 49 (original): The method of claim 38, wherein said non-effervescent flavored suspension is administered as part of a multisubstance regimen.

Claim 50 (original): The method of claim 49, wherein the color of the suspension identifies the biologically active substance to improve patient compliance with the multi-substance regimen.

Claim 51 (original): The method of claim 38, wherein said noneffervescent flavored suspension is administered to improve patient compliance with taking the biologically active substance.

Claim 52 (original): The method of claim 38, wherein the noneffervescent flavored suspension is administered to improve convenience of administration of the biologically active substance.

Claim 53 (original): The method of claim 38, wherein the solid dispersible tablet further contains a natural or artificial sweetening agent.

Claim 54 (original): The method of claim 38, wherein the biologically active substance is selected from the group consisting of analgesics, anti-inflammatories, antihistamines, antitussives, expectorants, decongestants, narcotics, bronchodilators, cardiovasculars, central nervous system drugs, anti-hypertensive agents, osteoporotic agents, GERD agents, anti-neoplastic agents, anti-asthmatics, hormone replacement agents, anti-infectives, antidiabetics, lipid lowering agents, thrombolytic agents, anticoagulant agents, fibrinolytic agents, nutritional agents, vitamins, minerals, metal salts, electrolytes, herbal agents and fatty acids.

Claim 55 (original): The method of claim 38, wherein the biologically active substance is an alkaline salt of potassium.

Claim 56 (original): The method of claim 55, wherein the alkaline salt of potassium is potassium chloride.

Claim 57 (original): The method of claim 38, wherein the liquid is water.

Claim 58 (original): The method of claim 38, wherein the non-effervescent flavored suspension has a pleasing taste when administered to the animal.

Claim 59 (original): The method of claim 38, wherein the non-effervescent flavored suspension is administered once a day.

Claim 60 (original): The method of claim 38, wherein the non-effervescent flavored suspension is administered at least twice a day.

Claims 61-65 (canceled)

Claim 66 (previously presented): The method of claim 38, wherein the animal is a mammal.

Claim 67 (currently amended): A method of improving patient compliance with a therapeutic regimen, which comprises:

administering to an animal a non-effervescent flavored suspension formed by placing into a liquid a solid dispersible tastemasked tablet comprising a flavoring agent and a plurality of particles being coated with an extended release coating agent;

wherein the solid dispersible tablet forms a non-effervescent flavored suspension when placed in a liquid having a viscosity of about 25 cp to about 75 cp; and

wherein the non-effervescent flavored suspension after being orally administered to the animal releases the biologically active substance over a period of about 2 hours to about 48 hours.

Claim 68 (previously presented): The method of claim 38, wherein the biologically active substance is a cardiovascular agent.